WE CLAIM:

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- 1. An aqueous nasal spray composition comprising a therapeutically effective amount of a medicament and an aqueous carrier comprising from 0.50 to 15.00% by weight/volume of a water soluble polymer selected from the group consisting of polyvinylpyrrolidone and mixtures thereof.
- 2. An aqueous nasal spray composition comprising of an effective amount of a medicament in an aqueous carrier 10 comprising:
 - 0.50 to 15.00% by weight/volume of a water soluble polymer selected from the group consisting of polyvinylpyrrolidone and mixtures thereof;
- 0.00 to 15.00% by weight/volume of polyethylene glycol;
 - 0.00 to 10.00% by weight/volume of a moisturizing agent or mixtures of moisturizing agents;
 - 0.00 to 10.00% by weight/volume of an antioxidant;
 - 0.001 to 0.10% by weight/volume of an antimicrobial preservative;
 - 2 0.00 to 5.00% by weight/volume of an aromatic alcohol;
- a sufficient amount of a pharmaceutically acceptable buffer to maintain the pH of the composition within the range of about 4.0 to 8.0 and

QS water.

3. The aqueous nasal spray composition of claim 2 comprising of an effective amount of a medicament in an aqueous carrier comprising:

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	0.50 to 2.5% by weight/volume of a
	water soluble polymer selected from the
	group consisting of polyvinylpyrrolidone
	and mixtures thereof;
5	0.5 to 10.00% by weight/volume of polyethylene glycol;
	1.00 to 4.00% by weight/volume of a moisturizing agent;
	0.01 to 0.05% by weight/volume of an
10	antioxidant:
	0.02 to 0.025% by weight/volume of an
	antimicrobial preservative;
	0.20 to 3.00% by weight/volume of an aromatic
	alcohol;
15	a sufficient amount of a pharmaceutically
	acceptable buffer to maintain the pH of the
	composition within the range of about 4.0 to
	8.0 and
	QS water.
20	Co manufacture and the control of th
_ •	4. The aqueous pasal spray composition of claim 2
	and added institution of claim 2
	comprising of an effective amount of a medicament in an
	aqueous carrier comprising:
٥.	1.00 to 1.50% by weight/volume of a
25	water soluble polymer selected from the
	group consisting of polyvinylpyrrolidone
	and mixtures thereof;
	2.5 to 5.00% by weight/volume of polyethylene
	glycol;
30	1.50 to 3.50% by weight/volume of a moisturizing
	agent or mixtures of moisturizing agents;
	0.015 to 0.030% by weight/volume of an
	antioxidant;

- 0.02 to 0.025% by weight/volume of an antimicrobial preservative;
- 0.25 to 1.00% by weight/volume of an aromatic alcohol:

a sufficient amount of a pharmaceutically acceptable buffer to maintain the pH of the composition within the range of about 4.0 to 8.0 and

QS water.

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- 5. The aqueous nasal spray composition of claim 2 wherein the moisturizing agent is propylene glycol; the antioxident is disodium EDTA; the antimicrobial preservative is benzalkonium chloride; and the aromatic alcohol is benzyl alcohol; and the buffer is a phosphate buffer.
- 6. The aqueous nasal spray composition of claim 5 wherein the medicament is selected from the group consisting of chlorpheniramine maleate, oxymetazoline hydrochloride 20 and mixtures thereof.
 - 7. The aqueous nasal spray composition of claim 6 wherein; the medicament is oxymetazoline hydrochloride.
- 25 8. The aqueous nasal spray composition of claim 6 wherein the medicament is chlorpheniramine maleate.
 - 9. The aqueous nasal spray composition of claim 2 which comprises:

30	<u>INGREDIENTS</u>	% Wt/Vol
	Water	QS
	Disodium EDTA	0.0200
	Sodium Phosphate Dibasic	0.0975
	Sodium Phosphate Monobasic	0.5525

	PVP K-90	0.2500
	PVP K-30	1.0000
	PEG 1450	2.5000
	Benzyl Alcohol	0.2500
5	Benzalkonium Chloride (17% solution)	0.0200
	Chlorpheniramine Maleate	0.5000
	Oxymetazoline Hydrochloride	0.0500

10. The aqueous nasal spray composition of claim 2 10 which comprises:

	INGREDIENTS	% Wt/Vol
	Water	QS
	Disodium EDTA	0.0200
1 5	Sodium Phosphate Dibasic	0.0975
	Sodium Phosphate Monobasic	0.5525
	PVP K-90	0.2500
	PVP K-30	1.0000
	PEG 1450	2.5000
20	Benzyl Alcohol	0.2500
	Benzalkonium Chloride (17% solution)	0.0200
	Oxymetazoline Hydrochloride	0.0500
	*	

11. The aqueous nasal spray compositions of claim 2 which comprises:

	<u>INGREDIENTS</u>	% Wt/Vol
	Water	QS
	Disodium EDTA	0.0200
3 0	Sodium Phosphate Dibasic	0.0975
	Sodium Phosphate Monobasic	0.5525
	PVP K-30	3.0000
	PEG 600	5.0000
	Benzyl Alcohol	0.2500

Benzalkonium Chloride (17% solution)	0.0200
Oxymetazoline Hydrochloride	0.0500
Chlorpheniramine Maleate	0.5000

5 12. The aqueous nasal spray composition of claim 2 which comprises:

	INGREDIENTS	% Wt/Vol
	Water	QS
10	Disodium EDTA	0.0200
	Sodium Phosphate Dibasic	0.0975
	Sodium Phosphate Monobasic	0.5525
	PVP K-30	3.0000
	PEG 1450	5.0000
15	Benzyl Alcohol	0.2500
	Benzalkonium Chloride (17% solution)	0.0200
	Oxymetazoline Hydrochloride	0.0500
	Chlorpheniramine Maleate	0.5000

20 13. The aqueous nasal spray composition of claim 2 which comprises:

	<u>INGREDIENTS</u>	% Wt/Vol
*	₹ Water	QS
25	Disodium EDTA	0.0200
	Sodium Phosphate Dibasic	0.0975
	Sodium Phosphate—Monobasic	0.5525
	PVP K-90	0.1000
	PVP K-30	3.0000
30	PEG 1450	2.5000
	Propylene glycol	0.2500
	Benzalkonium Chloride (17% solution)	0.1471
	Oxymetazoline Hydrochloride	0.0500

14. The aqueous nasal spray composition of claim 2 which comprises:

	<u>INGREDIENTS</u>	% Wt/Vol
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	Water	QS
	Disodium EDTA	0.0200
	Sodium Phosphate Dibasic	0.0975
	Sodium Phosphate Monobasic	0.5525
10	PVP K-90	0.1000
	PVP K-30	3.0000
	PEG 1450	5.0000
	Propylene Glycol	2.0000
	Glycerin	0.1000
15	Benzalkonium Chloride (17% solution)	0.1471
	Oxymetazoline Hydrochloride	0.0500

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